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AMENDMENT TO THE CLAIMS:

The following claim set replaces all prior versions, and listings, of claims in the application:

- (currently amended) Process for purification of a compound comprising an activated carbon treatment-using a filter unit containing activated carbon immobilized in a cartridge, the treatment comprising the steps of:
 - a) providing a filter unit containing activated carbon immobilized in a cartridge, wherein the cartridge is a self-contained replaceable entity comprising powdered activated carbon immobilized in a matrix which is in the form of a membrane sheet;
 - b) [[a)]] passing a first volume of a feed containing the compound over a first series of n connected filter units operating in series to obtain a first effluent, wherein n is at least two, said filter units having been assigned a position number 1 to n in the series and position number 1 being the first supplied with the feed.
 - c) [[b]] disconnecting a filter unit from the first series of filter units at any position number between 1 to n-1 after passing the first volume of feed, and connecting a fresh filter unit at any position that has a higher number than the position number of the disconnected filter unit, resulting in a next series of filter units,
 - (c))] passing a second volume of feed containing the compound over the next series of filter units to obtain a second effluent.
 - [d)]] optionally combining the effluents obtained in steps a) and c) to obtain a combined effluent, and
 - [e]]] recovering the compound from the first and second effluents, optionally the combined effluent.

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- (currently amended) The process according to claim 1, wherein step [[b]]] c)
 comprises disconnecting the filter unit at position number between 1 to n-1 and
 wherein the fresh filter unit is connected at position number n+1.
- (currently amended) The process according to claim 1, wherein step [[b)]] c)
 comprises disconnecting the filter unit at position number 1 and wherein the fresh
 filter unit is connected at position number n+1.
- (previously presented) The process according to claim 1 wherein the number n of connected filter units operating in series is 2 to 10.
- (previously presented) The process according to claim 1, wherein the treatment is operated in batch, semi-continuous or continuous mode.
- (previously presented) The process according to claim 1, wherein the flow rate of the feed is 0.05 to 400 L/min.
- (previously presented) The process according to claim 1 wherein the cartridge is a self-contained replaceable entity comprising powdered activated carbon immobilized in a matrix which is in the form of a membrane sheet.
- (previously presented) The process according to claim 7, wherein the flux over the membrane sheet is 1 to 50 L/m2/min.
- (previously presented) The process according to claim 1, wherein the residence time of the feed containing the compound in a single filter unit is at least 15 seconds and maximal 60 minutes.
- (previously presented) The process according to claim 1, wherein the process is operated at a temperature between minus 10°C to 40°C.

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 (previously presented) The process according to claim 1, wherein at least one disconnected filter unit is regenerated in situ by rinsing with a solvent.

12. (canceled)

- (previously presented) The process according to claim 1, wherein the compound is a secondary metabolite or a protein.
- (original) The process according to claim 13, wherein the secondary metabolite is selected from the group consisting of an antibiotic, a vitamin, a carotenoid or a PUFA.
- (previously presented) The process according to claim 1, wherein the compound is obtained by fermentation using a microorganism.
- (original) The process according to claim 14, wherein the microorganism is a Streptomyces species.
- (original) The process according to claim 15, wherein the Streptomyces species is selected from the group consisting of S. clavuligerus, S. coelicolor, S. griseus, S. venezuela, S. jumonjinensis, S. katsurahamanus or S. aureofaciens.
- (previously presented) The process according to claim 14, wherein the compound is selected from the group consisting of clavulanic acid, streptomycin, chloramphenicol, tetracycline or β-carotene.
- (previously presented) The process according to claim 1, further comprising the step of converting the compound into a pharmaceutically acceptable salt or food grade product.